

(19) World Intellectual Property Organization
International Bureau



(43) International Publication Date
15 March 2001 (15.03.2001)

PCT

(10) International Publication Number
WO 01/17577 A1

(51) International Patent Classification⁷: A61L 31/14,
31/16

(74) Agent: JOSEPHSON, Daryl, C.; Squire, Sanders
& Dempsey LLP, 600 Hansen Way, Palo Alto, CA
94304-1043 (US).

(21) International Application Number: PCT/US00/23535

(22) International Filing Date: 23 August 2000 (23.08.2000)

(25) Filing Language: English

(26) Publication Language: English

(30) Priority Data:
09/390,855 3 September 1999 (03.09.1999) US
09/390,069 3 September 1999 (03.09.1999) US

(71) Applicant: ADVANCED CARDIOVASCULAR SYS-
TEMS, INC. [US/US]; 3200 Lakeside Drive, Santa Clara,
CA 95054-2807 (US).

(72) Inventors: HOSSAINY, Syed, F., A.; 34325 Tupelo
Street, Fremont, CA 94555 (US). CHEN, Li; 5219 Iron-
shoe Drive, San Jose, CA 95138 (US).

(81) Designated States (*national*): AE, AL, AM, AT, AU, AZ,
BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK,
DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL,
IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU,
LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT,
RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA,
UG, UZ, VN, YU, ZA, ZW.

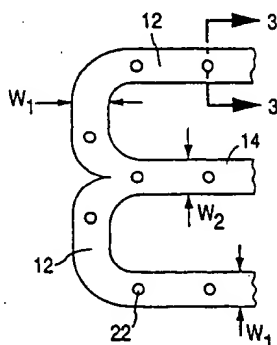
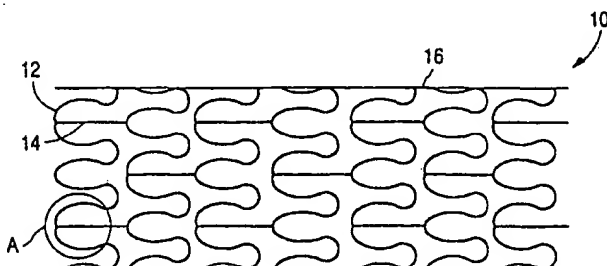
(84) Designated States (*regional*): ARIPO patent (GH, GM,
KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW), Eurasian
patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European
patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE,
IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG,
CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).

Published:

- With international search report.
- Before the expiration of the time limit for amending the
claims and to be republished in the event of receipt of
amendments.

[Continued on next page]

(54) Title: A POROUS PROSTHESIS AND A METHOD OF DEPOSITING SUBSTANCES INTO THE PORES



(57) Abstract: An implantable prosthesis, for example a stent, is provided that is capable of being loaded with substances for subsequent application to biological tissues. In one example, the prosthesis is a cylindrical-shaped body having depots or pores formed thereon. The depots can be formed at preselected locations on the body of the prosthesis and can have a preselected depth, size, and shape. The depots can have various shapes including a cylindrical or a conical shape. Substances such as therapeutic substances, polymeric materials, polymeric materials containing therapeutic substances, radioactive isotopes, and radiopaque materials can be deposited into the depots. A method of loading a substance into the prosthesis having depots or pores is also provided. A first fluid in combination with an added substance is applied to the porous prosthesis. During the application, the first fluid containing the substance is capable of penetrating into prosthesis pores. The first fluid is removed and a second fluid is applied to the prosthesis. The second fluid is not capable of significantly penetrating into the pores. Prior to the application of the second fluid, the prosthesis can be immersed in a third fluid and agitated via mechanical perturbation techniques so that any of the substance gathered on the surface of the body, after the application of the first fluid, is removed. The third

fluid should not be capable of dissolving the substance.

WO 01/17577 A1

WO 01/17577 A1



For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

**A POROUS PROSTHESIS
AND A METHOD OF DEPOSITING SUBSTANCES INTO THE PORES**

5

BACKGROUND OF THE INVENTION

Field of the Invention

This invention relates to implantable devices, such as an expandable,
10 intraluminal prosthesis commonly known as a stent. More particularly, this
invention relates to a prosthesis having pores formed in its cylindrical body.
Moreover, the present invention relates to a method of depositing substances, such
as therapeutic substances, in the pores.

Description of the Background

15 Percutaneous transluminal coronary angioplasty (PTCA) is a procedure for
treating heart disease. A catheter assembly having a balloon portion is introduced
into the cardiovascular system of a patient via the brachial or femoral artery. The
catheter assembly is advanced through the coronary vasculature until the balloon
portion is positioned across the occlusive lesion. Once in position across the
20 lesion, the balloon is inflated to a predetermined size to radially compress the
atherosclerotic plaque of the lesion against the inner wall of the artery to dilate the
lumen. The balloon is then deflated to a smaller profile to allow the catheter to be
withdrawn from the patient's vasculature.

A problem associated with the procedure includes formation of intimal
25 flaps or torn arterial linings which can collapse and occlude the conduit after the
balloon is deflated. Moreover, thrombosis and restenosis of the artery may
develop over several months after the procedure, which may require another
angioplasty procedure or a surgical by-pass operation. To reduce the partial or

total occlusion of the artery by the collapse of arterial lining and to reduce the chance of the development of thrombosis and restenosis, an intraluminal prosthesis, an example of which includes an expandable stent, is implanted in the lumen to maintain the vascular patency. Stents are scaffoldings, usually cylindrical or tubular in shape, functioning to physically hold open, and if desired, to expand the wall of the passageway. Typically stents are capable of being compressed for insertion through small cavities via small catheters, and then expanded to a larger diameter once at the desired location. Examples in patent literature disclosing stents which have been successfully applied in PTCA procedures include stents illustrated in U.S. Patent No. 4,733,665 issued to Palmaz, U.S. Patent No. 4,800,882 issued to Gianturco, and U.S. Patent No. 4,886,062 issued to Wiktor.

In treating the damaged vasculature tissue and to further fight against thrombosis and restenosis, there is a need for administering therapeutic substances to the treatment site. For example, anticoagulants, antiplatelets and cytostatic agents are commonly used to prevent thrombosis of the coronary lumen, to inhibit development of restenosis, and to reduce post-angioplasty proliferation of the vascular tissue, respectively. To provide an efficacious concentration to the treated site, systemic administration of such medication often produces adverse or toxic side effects for the patient. Local delivery is a highly suitable method of treatment in that smaller levels of medication, as compared to systemic dosages, are concentrated at a specific site. Local delivery produces fewer side effects and achieves more effective results.

One commonly applied technique for the local delivery of the drugs is through the use of medicated stents. One proposed method of medicating stents is to seed the stent with endothelial cells (Dichek, D.A. et al. Seeding of Intravascular Stents With Genetically Engineered Endothelial Cells; Circulation 1989; 80: 1347-1353). Briefly, endothelial cells can be seeded onto stainless steel stents and grown until the stents are covered. The cells are therefore able to be delivered to the vascular wall to provide therapeutic proteins. Another proposed method of providing therapeutic substances to the vascular wall includes simple heparin-

coated metallic stent, whereby a heparin coating is ionically or covalently bonded to the stent. Disadvantages associated with the aforementioned methods include significant loss of the therapeutic substance from the body of the stent during delivery and expansion of the stent and an absolute lack of control of the release rate of the therapeutic substance from the stent. Another proposed method involves the use of a polymeric carrier coated onto the body of the stent, as disclosed in U.S. Patent No. 5,464,650 issued to Berg et al., U.S. Patent No. 5,605,696 issued to Eury et al., U.S. Patent No. 5,865,814 issued to Tuch, and U.S. Patent No. 5,700,286 issued to Tartaglia et al. Obstacles often encountered with the use of a polymeric coating include difficulties in coating a complicated geometrical structure, poor adhesion of the polymeric coating to the surface of a stent, and biocompatibility of the polymer. Accordingly, it is desirable to be able to secure the therapeutic substance directly onto the body of the stent. Notwithstanding the benefits gained by securing a therapeutic substance to the body of the stent, it is also desirable to be able to secure other substances to the body of the stent, such as radiopaque materials, used to assist a physician to guide and deploy the stent at the proper site of treatment.

SUMMARY OF THE INVENTION

In accordance with various aspects of the present invention, an implantable prosthesis, one example of which includes a stent, is provided that is capable of being loaded with substances. The prosthesis is defined by a cylindrical shaped body having a thickness. Depots or pores are formed on the body at preselected locations. The depots have a preselected depth and shape. The depth of the depots can be equal to about 10% to about 90% of the thickness. In one embodiment, the depots can have a cylindrical shape. In another embodiment, the shape can be generally conical. Substances such as therapeutic substances, polymeric material, polymeric material containing therapeutic substances, radioactive isotopes, and radiopaque material can be deposited into the depots.

Another aspect of the present invention is a method of loading a substance into the depots. The method is applicable not only to the above-described

prosthesis, but to any type of porous prosthesis. A first fluid having a substance added therein is applied to a porous prosthesis. During the application, the first fluid containing the substance is capable of penetrating into the pores. The first fluid is removed, for example by evaporation, and a second fluid is applied to the
5 prosthesis. During the application of the second fluid, the second fluid is not capable of significantly penetrate into the pores. The second fluid can have a contact angle greater than about 90° . Contact angle is defied as the angle at the tangent of the fluid phase that has taken an equilibrium shape on a solid surface. In one embodiment of the present invention, the second fluid rinses the substance
10 from the surface of the body of the prosthesis. In another embodiment, a therapeutic substance and/or a polymer can be added to the second fluid to form a coating of a therapeutic substance and/or polymer onto the surface of the body of the prosthesis.

In accordance to another embodiment, prior to the application of the second
15 fluid, the prosthesis can be immersed in a third fluid and agitated via mechanical perturbation techniques. Accordingly, any of the substance gathered on the surface of the body after the application of the first fluid is removed. The third fluid should not be capable of dissolving the substance. The third fluid can have a contact angle above 90° .

20 BRIEF DESCRIPTION OF THE DRAWINGS

The features of the described embodiments are specifically set forth in the appended claims. However, embodiments relating to both structure and method of operation are best understood by referring to the following description and accompanying drawings, in which similar parts are identified by like reference
25 numerals.

Figure 1 is a side view of a conventional intraluminal prosthesis, the body of the prosthesis being defined by cylindrical elements engaged to one another by connecting elements;

Figure 2 is an enlarged view of section A of Figure 1, illustrating a portion of the cylindrical elements and connecting elements;

Figure 3A is a cross sectional view of the cylindrical element, taken in the direction of the arrows and along the plane of line 3-3 of Figure 2, illustrating a depot formed in the body of the prosthesis in accordance to one embodiment of the present invention;

Figure 3B is a cross sectional view of the cylindrical element, taken in the direction of the arrow and along the plane of line 3-3 of Figure 2, illustrating a depot formed in the body of the prosthesis in accordance to another embodiment of the present invention;

Figure 4A illustrates a fluid on a solid surface having a contact angle Φ_1 ; and

Figure 4B illustrates a fluid on a solid surface having a contact angle Φ_2 .

DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENTS

Figure 1 illustrates an implantable prosthesis 10, one example of which includes a stent. Stents are scaffoldings, usually cylindrical or tubular in shape, that are inserted into a anatomical passageway and operate to physically hold open and, if desired, to expand the wall of a passageway. Stents are capable of being compressed for insertion through small cavities via balloon-catheters, positioned in a desired location, then expanded to a larger diameter.

In one example, illustrated in Figures 1 and 2, stent 10 includes a plurality of rigid but resiliently flexible thread elements 12 that are arranged in a sinusoid-like configuration that is connected to form a continuous ring or cylinder. The plurality of cylindrical thread elements 12 are radially expandable, disposed coaxially, and interconnected by connecting elements 14 that are disposed between adjacent cylindrical thread elements 12, leaving gaps or lateral openings between adjacent cylindrical thread elements 12. Although the thread elements 12 are illustratively shown in the form of cylinders or rings connected axially-displaced

in-parallel, other configurations, such as helices, coils, or braids, and other connections may be utilized. Thread elements 12 and connecting elements 14 define a tubular stent body 16 having a lumen contacting surface 18 and an inner surface 20 as shown in Figures 3A and 3B.

5 Thread elements 12 have any suitable width W_1 , typically in a range of widths W_1 from about 0.002 inches to about 0.006 inches. A common width W_1 is about 0.003 inches. Connecting elements 14 have any suitable width W_2 , typically in a ranges of widths W_2 from about 0.002 inches to about 0.008 inches. A
10 common width W_2 is about 0.005 inches. Additionally, thread elements 12 have any suitable thickness T , typically a thickness T in a range from about 0.002 inches to about 0.008 inches. A common thickness T is about 0.005 inches. Connecting elements 14 have any suitable thickness, typically in a range from about 0.002 inches to about 0.008 inches. A common connecting element 14 thickness is about 0.005 inches. A specific choice of width and thickness depends on the anatomy
15 and size of the target lumen. In other words, the size of the stent can vary according to intended procedure, anatomy, and usage.

In one embodiment, thread elements 12 and connecting elements 14 are typically fabricated from a metallic material or an alloy such as stainless steel (e.g., 316L), "MP35N," "MP20N," tantalum, nickel-titanium alloy (commercially
20 available as Nitinol™), platinum-iridium alloy, gold, magnesium, or combinations of alloys. "MP35N" and "MP20N" are trade names for alloys of cobalt, nickel, chromium and molybdenum available from standard Press Steel Co., Jenkintown, PA. "MP35N" has a nominal composition of 35% cobalt, 35% nickel, 20% chromium, and 10% molybdenum. "MP20N" has a nominal composition of 50%
25 cobalt, 20% nickel, 20% chromium, and 10% molybdenum. The aforementioned list is merely a useful list of materials and that other materials are proven to function effectively.

A single depot or pore 22 or plurality of depots or pores 22 are formed as a laser trench or laser trenches on body 16 of stent 10 by exposing surface 18 to an
30 energy discharge from a laser, such as an excimer laser. Alternative methods of

forming depots 22 include physical or chemical etching techniques. Techniques of laser fabrication or etching to form depots 22 are well-known to one of ordinary skill in the art. Depots 22 can be formed in virtually any stent structure and not merely the above-described structure. The depots 22 are used for carrying a variety of substances including but not limited to therapeutic substances, polymers impregnated with therapeutic substances, radioactive isotopes, and radiopaque materials. The location of a depot 22 or depots 22 vary according to intended usage and application. Depots 22 are formed by a manufacturer at any preselected location and have any preselected depth, size, and geometrical configuration. In one example, depots 22 are evenly distributed through body 16 and have an equal volume so that the tissue in contact with stent 10 receives an equal distribution of a therapeutic substance. Depth D_1 of a depot 22 typically is varied in proportion to the thickness T of body 16 as well as the clinical purpose and usage.

For a stent 10 that carries a therapeutic substance or a polymeric carrier impregnated with a therapeutic substance, a suitable depot or pore depth D_1 has a range from about 10% to about 90 % of thickness T . Typically a depth not greater than about 50% of thickness T is most suitable. The specific depth D_1 of depots 22 depends on the amount of therapeutic substance that is to be deposited in depots 22. In an example of a stent 10 that carries a radioactive isotope, depth D_1 is typically about 10% to about 80% of thickness T . A more specific suitable depth is not greater than about 30% of thickness T is suitable.

For a stent 10 that carries a radiopaque material, a suitable depot or pore depth D_1 has a range from about 10% to about 90% of thickness T . Typically a depth not greater than about 65% is most suitable. A depth D_1 greater than about 65% of the thickness may compromise the structural integrity and mechanical functionality of stent 10. However the upper limit of depth D_1 varies depending on the material characteristics such as the hardness of the body 16.

Depots 22 may be formed in a variety of selected geometrical shapes. Referring to Figure 3A, a depot 22A is a generally cylindrical shape. A diameter D_2 of cylindrical depot 22A typically has a range from about 10% to about 90% of

width W_1 or W_2 , although the diameter D_2 is usually not greater than about 80% of width W_1 or W_2 . The specific diameter D_2 depends on the application and purpose of the depots 22. The upper limit of diameter D_2 varies depending on the material characteristics such as the hardness of the body 16.

5 An alternative example of a depot 22B, illustrated in Figure 3B, is generally conical in shape. Conical shaped depot 22B has an open end 24 and a closed end 26. The open end 24 is the end that contacts a surface of a tissue. A diameter D_3 of conical shaped depot 22B is shown to decrease from closed end 26 to open end 24. The largest diameter D_3' at the closed end 26 of conical shaped
10 depot 22B has a range from about 10% to about 80% of width W_1 or W_2 . Generally, the largest diameter D_3' is not greater than about 70% of width W_1 or W_2 . The smallest diameter D_3'' at the open end 24 of conical shaped depot 22B has a range from about 5% to about 70% of width W_1 or W_2 . Generally, the
15 smallest diameter D_3'' is not greater than about 60% of width W_1 or W_2 . The reduced size of opening 24 of conical shaped depot 22B, as compared to that of the cylindrical shaped depot 22A, reduces the rate at which a therapeutic substance is released once the stent is implanted at the desired location of treatment. The
depots 22 can have a variety of other geometrical shapes, such as elongated trenches (not illustrated).

20 The depth D_1 and diameters D_2 and D_3 of the individual depots 22 formed on body 16 of stent 10 can vary relative to one another. In one example, the manufacturer selectively controls the volume of depots on different positions of the body 16, either selectively varying the volume or making the volume consistent throughout the body 16. For some applications, consistent depot 22 volume is
25 important for delivery of a therapeutic substance to insure that the substance is evenly distributed throughout stent 10 and results in consistent application of the therapeutic substance to the tissues in contact with surface 18 of stent 10.

 A factor for determining the size, geometry, and concentration of depots 22 is the overall porosity of stent 10. Porosity is the total volume of pores in the body
30 16 of stent 10 divided by the total volume of structural material of the stent 10.

Porosity determines the capacity of substance that can be loaded into a stent 10 of predetermined dimensions. High porosity can adversely affect the structural integrity, strength, and elasticity of the stent 10. Consequently, stent design includes consideration of a tradeoff between strength, on one hand, and stent profile and stent load capacity on the other hand.

Substances are deposited into the depots or pores 22 using several illustrative methods. The methods are applicable to the illustrative stents 10 described hereinbefore and also to any type of porous prosthesis. In some examples, the deposited substance is a therapeutic substance or agent such as antineoplastics, antiinflammatory substances, antiplatelets, anticoagulants, fibrinolytics, thrombin inhibitors, antimitotics, and antiproliferatives. Examples of antineoplastics include paclitaxel and docetaxel. Examples of antiplatelets, anticoagulants, fibrinolytics, and thrombin inhibitors include sodium heparin, low molecular weight heparin, hirudin, argatroban, forskolin, vapiprost, prostacyclin and prostacyclin analogues, dextran, D-phe-pro-arg-chloromethylketone (synthetic antithrombin), dipyridamole, glycoprotein IIb/IIIa platelet membrane receptor antibody, recombinant hirudin, thrombin inhibitor (available from Biogen), and 7E-3B® (an antiplatelet drug from Centocore). Examples of suitable antimitotic agents include methotrexate, azathioprine, vincristine, vinblastine, flurouracil, adriamycin, and mutamycin. Examples of suitable cytostatic or antiproliferative agents include angiopeptin (a somatostatin analogue from Ibsen), angiotensin converting enzyme inhibitors such as Captopril® (available from Squibb), Cilazapril® (available from Hofman-LaRoche), or Lisinopril® (available from Merck); calcium channel blockers (such as Nifedipine), colchicine, fibroblast growth factor (FGF) antagonists, fish oil (omega 3-fatty acid), histamine antagonist, Lovastatin® (an inhibitor of HMG-CoA reductase, a cholesterol lowering drug from Merck), monoclonal antibodies (such as PDGF receptors), nitroprusside, phosphodiesterase inhibitors, prostaglandin inhibitor (available from Glaxo), Seramin (a PDGF antagonist), serotonin blockers, steroids, thioprotease inhibitors, triazolopyrimidine (a PDGF antagonist), and nitric oxide. Other therapeutic substances or agents which may be appropriate include alpha-

interferon, genetically engineered epithelial cells, and dexamethasone. While the listed therapeutic substances or agents are well known for preventative and therapeutic utility, the substances are listed by way of example and are not meant to be limiting. Other therapeutic substances which are currently available or that
5 may be developed in the future are equally applicable. The treatment of patients using the above mentioned medicines is well-known to those of ordinary skill in the art.

In other embodiments, the therapeutic substance is a radioactive isotope for stent usage in radiotherapeutic procedures. Examples of radioactive isotopes
10 include, but are not limited to, phosphoric acid ($\text{H}_3\text{P}^{32}\text{O}_4$), palladium (Pd^{103}), cesium (Cs^{131}), and iodine (I^{125}).

A therapeutic substance is added to a first fluid or solvent. The therapeutic substance is dispersed throughout the first solvent so that it is in a true solution, saturated or supersaturated with the solvent or suspended in fine particles in the
15 first solvent. If the therapeutic substance is suspended in particles in the first solvent, the pore size and the diameter of the opening of the pores are to be sufficiently large in comparison to the size of the particles to facilitate loading and unloading of the stent. In one example, suitable pores have a pore size that is more than ten times the particle size of a suspended therapeutic substance and an
20 opening diameter that is more than five times the diameter of the particle size.

The first solvent can be virtually any solvent that is compatible with the therapeutic substance. A suitable first solvent typically has a high capillary permeation. Capillary permeation or wetting is the movement of fluid on a solid substrate driven by interfacial energetics. Capillary permeation is quantitated by a
25 contact angle, defined as the angle at the tangent of the first solvent droplet in fluid phase that has taken an equilibrium shape on a solid surface. A low contact angle means a higher wetting liquid. A suitably high capillary permeation corresponds to a contact angle less than about 90° .

For example, Figure 4A illustrates a fluid having a high capillary permeation that corresponds to a solvent contact angle Φ_1 less than about 90° . Figure 4B illustrates a fluid having a low capillary permeation that corresponds to a fluid contact angle Φ_2 greater than about 90° . The first solvent can have a viscosity not greater than about ten centipoise. A high capillary permeation and a viscosity not greater than about ten centipoise allows the first solvent to penetrate into the pores of the prosthesis more quickly, eliminating a requirement to apply the first solvent to the prosthesis for a prolonged period of time. The first solvent can be volatile, facilitating evaporation of the first solvent. Useful examples of some first solvent include, but are not limited to, acetone, ethanol, methanol, isopropanol, tetrahydrofuran, and ethyl acetate. The first solvent is applied to a porous prosthesis, for example by immersing or spraying the solvent in procedures that are well-known to one having ordinary skill in the art.

The first solvent is applied for a predetermined period of time, the specific time depending on the capillary permeation and viscosity of the first solvent, the volume of the pores, and the amount of substance to be deposited. Therapeutic parameters such as the concentration of the therapeutic substance in the solvent and dosages depend on the duration of local release, the cumulative amount of release, and desired rate of release. Correlations and interrelations between the therapeutic parameters are well-known to one having ordinary skill in the art and are simply calculated.

After applying the first solvent for a selected duration, the first solvent is removed from the prosthesis. In one example, the first solvent is removed by evaporation in ambient pressure, room temperature, and anhydrous atmosphere and/or by exposure to mild heat (e.g., 60°C) under a vacuum condition. The prosthesis typically has a clustered or gross formation of a therapeutic substance gathered on the body surface. The cluster is generally removed by immersing the prosthesis in a second fluid and agitating the prosthesis via mechanical perturbation techniques, such as vortexing or vigorous shaking. The second fluid is a non-solvent so that the therapeutic substance does not significantly dissolve in

the second fluid. The non-solvent can have a low capillary permeation or a contact angle greater than about 90° and a viscosity not less than about 0.5 centipoise so that the second fluid is not capable of significantly penetrating into the pores during the process of agitation. Examples of a second fluid include but are not
5 limited to saturated hydrocarbons or alkanes, such as hexane, heptane, and octane.

The prosthesis is rinsed in a third fluid. The third fluid is typically a solvent to facilitate dissolution of the therapeutic substance. The third fluid generally has a low capillary permeation, corresponding to a contact angle greater than about 90° . The third fluid has a viscosity of not less than about 1.0 centipoise
10 and is therefore incapable of significantly penetrating into the pores during the rinsing stage. The rinsing is conducted rapidly for example in a range from 1 second to about 15 seconds, the exact duration depending on the solubility of the therapeutic substance in the solvent. Extended duration of exposure of the third solvent to the prosthesis may lead to the penetration of the third solvent into the
15 pores.

The rinsing step is repeated, if desired, until all traces of therapeutic substance are removed from the surface of the stent. The third fluid removes excess traces of therapeutic substance from the surface of the prosthesis body. Useful examples of third fluids include but are not limited to dimethylsulfoxide
20 (DMSO), water, DMSO in an aqueous solution, glyme, and glycerol. The third fluid is removed from the prosthesis body using a technique such as evaporation in ambient pressure, room temperature and anhydrous atmosphere and/or by exposure to mild heat (e.g., 60°C) under vacuum condition. The first, second, third fluids are selected to not adversely affect the characteristics and composition of the
25 therapeutic substance.

In one embodiment, the third fluid can be highly volatile, for example having a boiling point of not greater than about 60°C at 1 atm. Accordingly, the third fluid is capable of rapidly evaporating. Rapid evaporation of the third fluid causes the third fluid to be removed from the prosthesis prior to any significant

penetration of the third fluid in the pores. A useful example of a highly volatile third fluid includes, but is not limited to, Freon (e.g., Xerosolv™).

Once loaded, the therapeutic substance remains in the pores until prosthesis deployment and expansion. The expanded prosthesis engages the wall of the anatomical passageway and the therapeutic substance disseminates from the porous cavities and is absorbed into the tissue of the walls of the passageway that are in contact with the prosthesis.

In some embodiments, a surface of the stent is coated with a therapeutic substance in addition to having a therapeutic substance deposited in the pores. A coating of therapeutic substance on the surface of the prosthesis is formed by adding the therapeutic substance to the third fluid rinse. The therapeutic substance is dispersed through the third fluid to form a true solution with the third solvent, rather than a dispersion of fine particles. The therapeutic substance is a substance that is capable of absorbing or attaching to the prosthesis surface. For example, highly suitable therapeutic substances for a stainless steel prosthesis include taxol and dexamethasone. Suitable substances for a Nitinol™ prosthesis include aspirin and heparin. The therapeutic substance added to the third fluid can be the same substance as the therapeutic substance deposited in the pores or a different substance. Rinsing with the third fluid is optionally prolonged or repeated to increase the thickness of the prosthesis therapeutic substance coating.

In another example, a polymeric coating is formed on the surface of the prosthesis, covering the pores containing deposited therapeutic substance. The polymeric coating forms a membrane that reduces the rate of release of a therapeutic substance from the pores. A polymeric material is added to the third fluid rinse to form a coating made from the polymeric material on the prosthesis surface.

The polymeric material, by example and not limitation, forms about 1% to about 3% by weight of the total weight of the solution. The polymeric material is most suitably bio-compatible, including polymers that are non-toxic, non-

inflammatory, chemically inert, and substantially non-immunogenetic in the applied amounts. The polymer is typically either be bioabsorbable or biostable. A bioabsorbable polymer bio-degrades or breaks down in the body and is not present sufficiently long after implantation to cause an adverse local response.

- 5 Bioabsorbable polymers are gradually absorbed or eliminated by the body by hydrolysis, metabolic process, bulk, or surface erosion. Examples of bioabsorbable, biodegradable materials include but are not limited to polycaprolactone (PCL), poly-D, L-lactic acid (DL-PLA), poly-L-lactic acid (L-PLA), poly(lactide-co-glycolide), poly(hydroxybutyrate), poly(hydroxybutyrate-co-valerate), polydioxanone, polyorthoester, polyanhydride, poly(glycolic acid),
10 poly(glycolic acid-cotrimethylene carbonate), polyphosphoester, polyphosphoester urethane, poly (amino acids), cyanoacrylates, poly(trimethylene carbonate), poly(iminocarbonate), copoly(ether-esters), polyalkylene oxalates, polyphosphazenes, polyiminocarbonates, and aliphatic polycarbonates.
- 15 Biomolecules such as heparin, fibrin, fibrinogen, cellulose, starch, and collagen are typically also suitable. Examples of biostable polymers include Parylene®, Parylast®, polyurethane (for example, segmented polyurethanes such as Biospan®), polyethylene, polyethylene terephthalate, ethylene vinyl acetate, silicone and polyethylene oxide.

- 20 After the evaporation of the third solvent, a polymeric layer remains on the surface of the prosthesis and over the pores. The polymeric coating is alternatively formed by other conventional methods such as plasma polymerization, the practice of which is well known to one of ordinary skill in the art.

- In another example, the aforementioned polymeric coating is impregnated
25 with a therapeutic substance that is added to the third fluid-polymer mixture. Concentration of the therapeutic substance depends on release parameters including the duration of local release, the cumulative amount of release, and the desired rate of release. The correlations and interrelations between the release parameters are well-known to those of ordinary skill in the art and easily
30 calculated. The polymeric material, by example and not limitation, makes up from

about 1% to about 3% by weight of the total weight of the solution. The therapeutic substance, by example and not limitation, typically makes up about 0.3% to about 1% of the total weight of the solution. Once the third solvent is evaporated, a polymeric coating impregnated with a therapeutic substance remains
5 on the surface, covering the therapeutic substance-filled pores.

In another example, a polymeric material, such as any of the polymeric materials listed herein, is impregnated with a therapeutic substance and deposited into the pores. The polymeric material reduces the rate of release of the therapeutic substance from the pores. The method of application includes adding a
10 therapeutic substance and a polymer to a first fluid or solvent. The therapeutic substance is dispersed throughout the first solvent to dissolve into a true solution, or is saturated or supersaturated with the solvent or suspended in fine particles in the first solvent. The polymeric material is also dispersed throughout the first solvent to form a true solution, or is suspended in fine particles in the first solvent.
15 Saturation or supersaturation of the polymer is less suitable since the viscosity of the first solvent is raised beyond a desired limit.

If the therapeutic substance is suspended in the first solvent, the pore size and the diameter of the opening of the pores are to be sufficiently large in comparison to the size of the particles to facilitate loading and unloading of the
20 stent. In one example, suitable pores have a pore size that is more than ten times the particle size of a suspended therapeutic substance and an opening diameter that is more than five times the diameter of the particle size.

The first solvent is selected from among solvents that are compatible with the polymer and therapeutic substance. A first solvent having a high capillary
25 permeation or a contact angle not higher than about 90° improves performance. The first solvent can have a viscosity not higher than about ten centipoise. The first solvent can be volatile to facilitate evaporation. Useful examples of a first solvent include but are not limited to acetone, ethanol, methanol, isopropanol, tetrahydrofuran, and ethyl acetate. The first solvent is applied to a porous
30 prosthesis, by for example, immersing or spraying. The first solvent is applied for

a predetermined time period, the specific time depending on the capillary permeation and viscosity of the first solvent, volume of the pores, and the amount of substance to be deposited. Therapeutic parameters such as the concentration of the therapeutic substance and the specific polymer depend on the duration of local
5 release, the cumulative amount of release, and the rate of release that is desired. Correlations and interrelations between the therapeutic parameters are well-known to those having ordinary skill in the art and are easily determined.

After a selected time duration, the first solvent is removed from the prosthesis by a technique such as evaporation in ambient pressure, room
10 temperature, and anhydrous atmosphere and/or by exposure to mild heat (e.g., 60° C) under vacuum condition. The prosthesis typically has a cluster of polymeric material and therapeutic substance formed on the body surface. The cluster is removed by immersing the prosthesis in a second fluid and agitating the prosthesis via mechanical perturbation techniques, such as vortexing or vigorous shaking.
15 The second fluid is a non-solvent so that the therapeutic substance and the polymer are not capable of dissolution in the second fluid. The non-solvent generally has a low capillary permeation, corresponding to a contact angle greater than about 90° and a viscosity not less than 0.5 centipoise so that the second fluid is not capable of significantly penetrating into the pores during the process of agitation. Examples
20 of the second fluid include but are not limited to saturated hydrocarbons or alkanes, such as hexane, heptane, and octane.

The prosthesis is rinsed in a third fluid subsequent to the application of the first solvent or the agitation of the prosthesis. The third fluid typically has a low capillary permeation, corresponding to a contact angle greater than about 90°. The
25 third fluid has a viscosity of not less than 1.0 centipoise. The third fluid is not capable of significantly penetrating into the pores during the rinsing stage. In one embodiment, the third fluid is capable of significantly dissolving both the therapeutic substance and the polymeric material. Rinsing is conducted rapidly for example for a duration in a range from 1 second to about 15 seconds. The specific
30 duration depends on the solubility of the therapeutic substance and the polymeric

material characteristics. The rinsing is repeated, if desired, until all traces of therapeutic substance and polymeric material are removed from the stent surface.

In an alternative embodiment, the third fluid is capable of significantly dissolving the therapeutic substance but not the polymeric material. As a result, traces of therapeutic substance are removed from the surface of the prosthesis, leaving a polymeric coating covering the surface of the prosthesis including the pores. The polymeric coating serves as an additional rate-reducing membrane. Useful examples of third fluid include but are not limited to DMSO, water, DMSO in an aqueous solution, and glycerol. The third fluid can be removed from the body of the prosthesis using a technique such as evaporation in ambient pressure, room temperature and anhydrous atmosphere and/or by exposure to mild heat (e.g., 60° C) under vacuum condition. The first, second, third fluids, alone or in conjunction with the polymeric material should not adversely affect the characteristics and composition of the therapeutic substance.

In another example, a polymeric material, such as a material capable of swell-loading or post-loading, can be deposited into the pores. Swell-loading occurs when a polymeric carrier is soaked with a therapeutic substance/solvent solution. The polymer swells, receiving the therapeutic substance in the polymer matrices. Once the solvent is removed, the polymer collapses and is impregnated with the therapeutic substance. Examples of polymeric material that are susceptible to swell-loading include thermoplastic polymers such as polyurethanes, polylactic acid, and polyglycolic acid, and non-thermoplastic polymers such as polyethyleneglycol, polyvinyl alcohol, polyacrylamide, and tecophilic polymers. The method of depositing a polymeric material into the pores is generally similar to the above-described methods with an addition of a curing step for the non-thermoplastic polymers subsequent to the rinsing step. One of ordinary skill in the art of polymer fabrication understands how to cure a non-thermoplastic polymer.

In another example, a radiopaque substance such as gold is deposited into the pores. The process of depositing radioactive isotopes is generally similar to the methods described above with a radiopaque substance dispersed and suspended in

fine particles through a first fluid. The first fluid can have a high capillary permeation or a contact angle not higher than about 90°. The first fluid has a viscosity not higher than about ten centipoise and can be volatile, ensuring that the first fluid evaporates more readily and easily. Useful examples of first fluids
5 include acetone, ethanol, methanol, isopropanol, tetrahydrofuran, and ethyl acetate.

The first fluid is applied to a porous prosthesis for a predetermined period of time, typically about 30 minutes. After application, the first solvent is removed from the prosthesis using a technique such as evaporation in ambient pressure, room temperature, and anhydrous atmosphere and/or by exposure to mild heat (for
10 example, 60° C) under vacuum condition. The prosthesis may have a gross formation of radiopaque substance gathered on the prosthesis body surface. The radiopaque formation can be removed by immersing the prosthesis in a second fluid and agitating the prosthesis via mechanical perturbation techniques, such as vortexing or vigorous shaking. The second fluid can have a low capillary
15 permeation or a contact angle greater than about 90° and a viscosity not less than about 0.5 centipoise so that the second fluid is not capable of significantly penetrating into the pores during the process of agitation. Examples of the second fluid include but are not limited to saturated hydrocarbons or alkanes, such as hexane, heptane, and octane.

20 The prosthesis is rinsed in a third fluid subsequent to the application of the first fluid or the agitation of the prosthesis. The third fluid has a low capillary permeation or a contact angle greater than about 90°, and has a viscosity not less than about 1.0 centipoise so that the third fluid is not capable of significantly penetrating into the pores during the rinsing stage. The rinsing is conducted
25 rapidly, for example in a range from 1 second to about 15 seconds, since an extended exposure duration may result in penetration of the third fluid into the pores. Suitable materials for the third fluid include DMSO, water, glyme and glycerol. Rinsing is repeated, if desired, until all traces of radiopaque substance are removed from the surface of the stent. The third fluid is removed from the
30 body of the prosthesis using a technique such as evaporation in ambient pressure,

room temperature and anhydrous atmosphere or by exposure to mild heat (e.g., 60° C) under vacuum condition. Sintering of the radiopaque material deposited in the pores is performed to bond particles of the radiopaque material without melting the particles. Appropriate pressure and temperature of radiopaque material sintering is
5 specific to the particular material in a manner well known to one having ordinary skill in the art.

Several examples illustrate various methods for depositing substances such as therapeutic substances on a stent. The examples illustrate but do not limit the possible techniques for depositing substances.

10

EXAMPLE 1

Trapidil is dissolved in ethanol by conventional methods. Trapidil makes up about 15% by weight of the total weight of the solution. A stent having a porous surface is immersed in the solution for 30 minutes. The stent is removed and mounted on a mandrel at ambient pressure, room temperature, and anhydrous
15 atmosphere for approximately 30 minutes, until the ethanol is evaporated. The stent is submerged in hexane, followed by mechanical perturbation in a vortex apparatus for about 15 seconds. The stent is removed from the non-solvent and rinsed with water for about 5 seconds. The stent is dried at ambient pressure, room temperature, and anhydrous atmosphere.

20

EXAMPLE 2

Trapidil is dissolved in ethanol by conventional methods. Trapidil makes up about 20% by weight of the total weight of the solution. A stent having a porous surface is immersed in the solution for 20 minutes. The stent is removed and mounted on a mandrel at ambient pressure, room temperature, and anhydrous
25 atmosphere for approximately 30 minutes, until the ethanol is evaporated. The stent is submerged in heptane, followed by mechanical perturbation in vortex apparatus for 45 seconds. The stent is removed from the non-solvent and rinsed with dimethylsulfoxide (DMSO) for about 5 seconds. After rinsing, the mandrel is

placed in ambient pressure, room temperature, and anhydrous atmosphere for approximately 2 hours, until the DMSO is evaporated.

EXAMPLE 3

Trapidil is dissolved in ethanol by conventional methods. Trapidil makes
5 up about 25% by weight of the total weight of the solution. A stent having a
porous surface is immersed in the solution for 20 minutes. The stent is removed
and mounted on a mandrel at ambient pressure, room temperature, and anhydrous
atmosphere for approximately 30 minutes, until the ethanol is evaporated. The
stent is submerged in heptane, followed by mechanical perturbation in vortex
10 apparatus for about 60. The stent is removed from the non-solvent and rinsed with
a dimethylsulfoxide (DMSO) aqueous solution (having a 1:1 DMSO-water ratio)
for about 5 seconds. After rinsing, the mandrel is placed in ambient pressure, room
temperature, and anhydrous atmosphere for approximately 2 hours. Next, the
prosthesis was placed in an oven under vacuum condition and at a temperature of
15 about 60° C for 24 hours, until all of the DMSO and water evaporated.

EXAMPLE 4

Trapidil is dissolved in ethanol by conventional methods. Trapidil makes
up about 25% by weight of the total weight of the solution. A stent having a
porous surface is immersed in the solution for 30 minutes. The stent is removed
20 and mounted on a mandrel at ambient pressure, room temperature, and anhydrous
conditions for approximately 30 minutes, until the ethanol is evaporated. The stent
is submerged in heptane, followed by mechanical perturbation in a vortex
apparatus for about 45 seconds. The stent is removed from the non-solvent and
rinsed, for about 10 seconds, with solution of ethylene vinyl alcohol and DMSO,
25 the ethylene vinyl alcohol constituting about 1% by weight of the total weight of
the solution. The mandrel is placed in ambient pressure, room temperature, and
anhydrous condition for approximately 2 hours. The stent is placed in a oven
under vacuum condition and at a temperature of about 60° C for 24 hours, until all

of the DMSO is evaporated. A polymeric coating remains on the surface of the stent.

EXAMPLE 5

Trapidil is dissolved in ethanol by conventional methods. Trapidil makes
5 up about 20% by weight of the total weight of the solution. A stent having a
porous surface is immersed in the solution for 40 minutes. The stent is removed
and mounted on a mandrel at ambient pressure, room temperature, and anhydrous
conditions for approximately 30 minutes, until the ethanol is evaporated. The stent
is submerged in heptane, followed by mechanical perturbation in a vortex
10 apparatus for about 30 seconds. The stent is removed from the non-solvent and
rinsed, for about 10 seconds, with a solution containing ethylene vinyl alcohol,
trapidil, and DMSO. The ethylene vinyl alcohol constitutes about 1% by weight
and the trapidil constitutes about 0.33% by weight of the total weight of the
solution. The mandrel is placed in ambient pressure, room temperature, and
15 anhydrous condition for approximately 2 hours. The stent is placed in a oven
under vacuum condition and at a temperature of about 60° C for 24 hours, until all
of the DMSO is evaporated. A polymeric coating having a trapidil impregnated
therein remains on the stent.

EXAMPLE 6

20 Trapidil was dissolved in ethanol by conventional methods. Trapidil was
used to make 40% by weight of the total weight of the solution. A stent having a
porous surface was mounted on a mandrel and dipped in the 40% solution for 40
seconds. The stent was removed from the other end and dried at ambient pressure,
room temperature, and anhydrous conditions for about 30 minutes, until the
25 ethanol was evaporated. The stent was mounted on a mandrel again and rinsed in a
heparin (DuraFlo™)/ Trapidil solution for 3 seconds. The solution constituted
0.6% by weight of Heparin and 0.6% by weight of Trapidil. The solvent used was
Freon (Xerosolv) and n-Propanol in the ratio of 5:1 by volume. The stent was
placed in a humidity controlled chamber at room temperature for 12 hours until all

of the Freon and n-Propanol solvent system evaporated. A DuraFlo coating having Trapidil impregnated therein remained on the stent surface and inside the pores.

While particular embodiments of the present invention have been shown and described, it will be obvious to those skilled in the art that changes and
5 modifications can be made without departing from this invention in its broader aspects and, therefore, the appended claims are to encompass within their scope all such changes and modifications as fall within the true spirit and scope of the invention.

CLAIMS

What is claimed is:

1. A method according to loading a substance into pores of an
5 implantable prosthesis:
providing a prosthesis having a surface and pores formed in said
surface;
applying a first fluid having an added substance to said prosthesis,
wherein during said act of applying, said first fluid penetrates into said
10 pores;
removing said first fluid from said prosthesis substantially to
elimination;
applying a second fluid to said prosthesis, wherein during said act
of applying said second fluid, said second fluid does not significantly
15 penetrate into said pores; and
removing said second fluid from said prosthesis substantially to
elimination, wherein said substance is deposited into said pores.
2. The method according to Claim 1, wherein said acts of removing
20 said first fluid and said second fluid comprise evaporating said first fluid and said
second fluid from said prosthesis.
3. The method according to Claim 1, additionally comprising prior to
said act of applying said second fluid:
25 immersing said prosthesis in a third fluid, wherein said substance
does not significantly dissolve in said third fluid;
agitating said prosthesis in said third fluid to significantly remove
said substance from said surface of said prosthesis, wherein said substance
remains in said pores; and
30 removing said prosthesis from said third fluid.

4. The method according to Claim 3, wherein said third fluid has a contact angle greater than about 90°.

5. The method according to Claim 1, wherein said substance
5 significantly dissolves when in contact with said second fluid.

6. The method according to Claim 1, wherein said second fluid has a contact angle greater than about 90°.

10 7. The method according to Claim 1, wherein said substance is a first therapeutic substance.

8. The method according to Claim 7, wherein said first therapeutic substance significantly dissolves when in contact with said second fluid.

15

9. The method according to Claim 7, wherein said second fluid has a contact angle greater than about 90°.

10. The method according to Claim 7, wherein said second fluid has a
20 boiling point of not greater than about 60°

11. The method according to Claim 7, additionally comprising, prior to said act of applying said second fluid:

immersing said prosthesis in a third fluid, wherein said first
25 therapeutic substance does not significantly dissolve in said third fluid;
agitating said prosthesis in said third fluid to significantly remove
said first therapeutic substance from said surface of said prosthesis, wherein
said first therapeutic substance remains in said pores; and
removing said prosthesis from said third fluid.

30

12. The method according to Claim 11, wherein said third fluid has a contact angle greater than about 90°.

13. The method according to Claim 7, wherein said second fluid comprises a second therapeutic substance added thereto, such that after said act of significantly removing said second fluid from said prosthesis, a coating of said second therapeutic substance remains on said surface of said prosthesis.

5

14. The method according to Claim 13, wherein said second therapeutic substance is the same as the first therapeutic substance.

15. The method according to Claim 13, wherein said second therapeutic
10 substance is different than said first therapeutic substance.

16. The method according to Claim 7, wherein said second fluid comprises a polymeric material added thereto, such that after said act of significantly removing said second fluid from said prosthesis, a coating of said
15 polymeric material remains on said surface of said prosthesis.

17. The method according to Claim 7, wherein said second fluid comprises a combination of a polymeric material and a second therapeutic substance added thereto, such that after said act of significantly removing said
20 second fluid from said prosthesis, a coating of said polymeric material containing said second therapeutic substance remains on said surface of said prosthesis.

18. The method according to Claim 1, wherein said substance is a polymeric material.

25

19. The method according to Claim 18, wherein said polymeric material significantly dissolves when in contact with said second fluid.

20. The method according to Claim 18, wherein said second fluid has a
30 contact angle greater than about 90°.

21. The method according to Claim 18, additionally comprising, prior to said act of applying said second fluid:

immersing said prosthesis in a third fluid, wherein said polymeric material does not significantly dissolve in said third fluid;

5 agitating said prosthesis in said third fluid to significantly remove said polymeric material from said surface of said prosthesis, wherein said polymeric material remains in said pores; and

removing said prosthesis from said third fluid.

10 22. The method according to Claim 21, wherein said third fluid has a contact angle greater than about 90°.

23. The method according to Claim 18, wherein said polymeric material is a non-thermoplastic polymer and the method additionally comprises the act of
15 curing said non-thermoplastic polymer deposited in said pores.

24. The method according to Claim 1, wherein said substance is a combination of a polymeric material and a therapeutic substance.

20 25. The method according to Claim 24, additionally comprising, prior to said act of applying said second fluid:

immersing said prosthesis in a third fluid, wherein said polymeric material and said therapeutic substance do not significantly dissolve in said third fluid;

25 agitating said prosthesis in said third fluid to significantly remove said polymeric material and said therapeutic substance from said surface of said prosthesis, wherein said polymeric material containing said therapeutic substance remains in said pores; and

removing said prosthesis from said third fluid.

30 26. The method according to Claim 25, wherein said third fluid has a contact angle greater than about 90°.

27. The method according to Claim 24, wherein said polymeric material and said therapeutic substance dissolves when in contact with said second fluid such that any of said polymeric material and said therapeutic substance disposed on said surface of said prosthesis are significantly removed.

5

28. The method according to Claim 24, wherein said therapeutic substance dissolves when in contact with said second fluid, wherein after said act of significantly removing said second fluid a coating made from said polymeric material remains on said surface of said prosthesis and covers said pores.

10

29. The method according to Claim 24, wherein said second fluid has a contact angle greater than about 90°.

30. The method according to Claim 1, wherein said substance is a radioactive isotope.

15

31. The method according to Claim 30, wherein said second fluid has a contact angle greater than about 90°.

20 32. The method according to Claim 30, additionally comprising prior to said act of applying said second fluid:

immersing said prosthesis in a third fluid;

agitating said prosthesis in said third fluid to significantly remove said radioactive isotope from said surface of said prosthesis; and

25 removing said prosthesis from said third fluid.

33. The method according to Claim 32, wherein said third fluid has a contact angle greater than about 90°.

30 34. The method according to Claim 1, wherein said substance is a radiopaque material.

35. The method according to Claim 34, wherein said second fluid has a contact angle greater than about 90°.

36. The method according to Claim 34, additionally comprising, subsequent to said act of significantly removing said second fluid, sintering said radiopaque material deposited in said pores.

37. The method according to Claim 34, additionally comprising prior to said act of applying said second fluid:

immersing said prosthesis in a third fluid;
agitating said prosthesis in said third fluid to significantly remove said radiopaque from said surface of said prosthesis; and
removing said prosthesis from said third fluid.

38. The method according to Claim 37, wherein said third fluid has a contact angle greater than about 90°.

39. A method according to depositing a substance into a plurality of pores formed in a body of an implantable prosthesis, comprising:

providing a first fluid having an added substance;
applying said first fluid to a prosthesis, said prosthesis having a body and a plurality of pores formed in said body, wherein said first fluid is applied until a predetermined amount of said substance has penetrated into said pores;

removing said first fluid from said prosthesis substantially to elimination wherein said added substance remains in said pores;

immersing said prosthesis in a second fluid, wherein said substance does not significantly dissolve in said second fluid;

agitating said prosthesis in said second fluid to significantly remove said substance from a surface of said body;

removing said prosthesis from said second fluid;

rinsing said prosthesis with a third fluid, wherein during said act of rinsing, said third fluid is not capable of significantly penetrating into said pores; and

significantly removing said third fluid from said prosthesis, wherein said substance is deposited in said pores.

40. The method according to Claim 39, wherein said first fluid has a contact angle less than about 90°.

41. The method according to Claim 39, wherein during said acts of
5 immersing and agitating said second fluid is not capable of significantly penetrate into said pores.

42. The method according to Claim 39, wherein said second fluid has a lower capillary permeation than said first fluid.

10 43. The method according to Claim 39, wherein said second fluid has a contact angle greater than about 90°.

44. The method according to Claim 39, wherein said substance
15 dissolves when in contact with said third fluid.

45. The method according to Claim 39, wherein said third fluid has a lower capillary permeation than said first fluid.

20 46. The method according to Claim 39, wherein said third fluid has a contact angle greater than about 90°.

47. An implantable prosthesis, comprising:

25 a body structure having a generally cylindrical shape, an outer surface capable of contacting an inner lumen surface of a passageway, and a hollow bore which extends longitudinally through said body structure to define an inner surface, said body structure having a thickness;

30 said body structure having a plurality of depots formed therein, said depots having a depth less than said thickness of said body structure, and said depots having a preselected and controlled distribution and a preselected and controlled depth.

48. The implantable prosthesis of Claim 47, wherein said prosthesis is a stent.

49. The implantable prosthesis of Claim 47, wherein said depots are
5 laser trenches formed by exposing said surface to an energy discharge from a laser.

50. The implantable prosthesis of Claim 47, wherein said preselected and controlled depth is equal to about 10% to 90% of said thickness.

10 51. The implantable prosthesis of Claim 47, wherein said preselected and controlled depth is not greater than about 65% of said thickness.

52. The implantable prosthesis of Claim 47, wherein said depots have a preselected shape.

15

53. The implantable prosthesis of Claim 52, wherein said preselected depot shape is generally cylindrical.

54. The implantable prosthesis of Claim 52, wherein said preselected
20 depot shape is generally conical.

55. The implantable prosthesis of Claim 54, wherein said conical shaped depot has an open end and a closed end, said open end contacts said passageway inner lumen surface, and said open end has a diameter smaller than a
25 diameter of said closed end.

56. The implantable prosthesis of Claim 47 further comprising a substance deposited into said depots.

30 57. The implantable prosthesis of Claim 47 further comprising a therapeutic substance deposited into said depots.

58. The implantable prosthesis of Claim 47 further comprising a therapeutic substance deposited into said depots, wherein said therapeutic substance is selected from a group of antineoplastic, antiplatelet, anticoagulant, fibrinolytics, antimitotic, thrombin inhibitor, antiinflammatory, and antiproliferative substances.

59. The implantable prosthesis of Claim 47 further comprising a therapeutic substance deposited into said depots, wherein said therapeutic substance is a radioactive isotope.

60. The implantable prosthesis of Claim 47 further comprising a substance deposited into said depots, wherein said substance is a polymeric material.

61. The implantable prosthesis of Claim 47 further comprising a substance deposited into said depots, wherein said substance is a polymeric material containing a therapeutic substance.

62. The implantable prosthesis of Claim 47 further comprising a substance deposited into said depots, wherein said substance is a polymeric material containing a therapeutic substance, said therapeutic substance is selected from a group of antineoplastic, antiplatelet, anticoagulant, fibrinolytic, antimitotic, thrombin inhibitor, antiinflammatory, and antiproliferative substances.

63. The implantable prosthesis of Claim 47 further comprising a substance deposited into said depots, wherein said substance is a radiopaque substance.

64. The implantable prosthesis of Claim 47 further comprising a first therapeutic substance deposited in said depots and a polymeric coating formed on said lumen contacting surface of said body, wherein said polymeric coating covers said depots and reduces the rate at which said first therapeutic substance is released from said depots.

65. The implantable prosthesis of Claim 64, wherein said polymeric coating contains a second therapeutic substance.

66. A method of loading a substance into a body of a implantable
5 prosthesis, comprising:

forming depots in preselected locations of said implantable
prosthesis body; and

depositing a substance into said depots.

67. The method according to Claim 66, further comprising:
10 forming a body structure having a generally cylindrical shape, an
outer surface capable of contacting an inner lumen surface of a
passageway, and a hollow bore which extends longitudinally through said
body structure to define an inner surface, said body structure having a
thickness.

15

68. The method according to Claim 66 further comprising:
controlling formation of the depots to a predetermined depth.

69. The method according to Claim 66 further comprising:
20 controlling formation of the depots to a predetermined geometrical
shape.

1/1

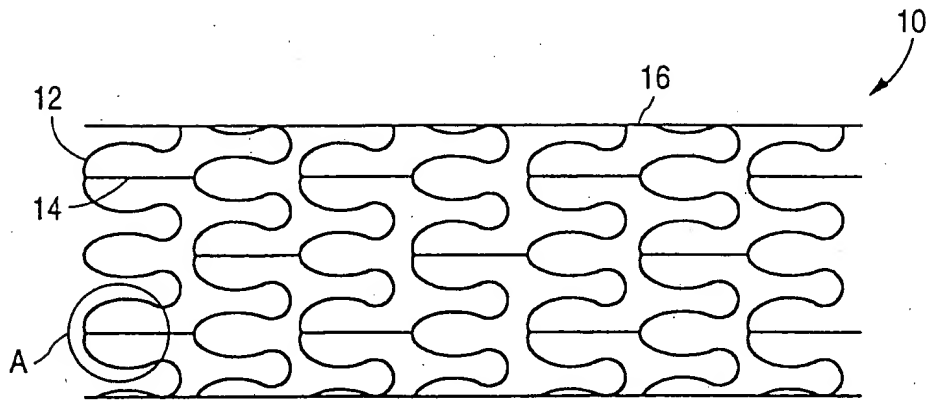


Fig. 1

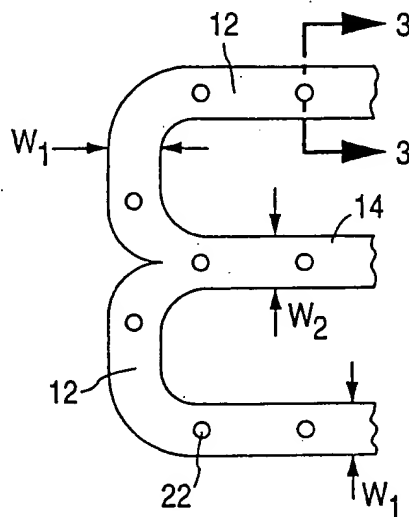


Fig. 2

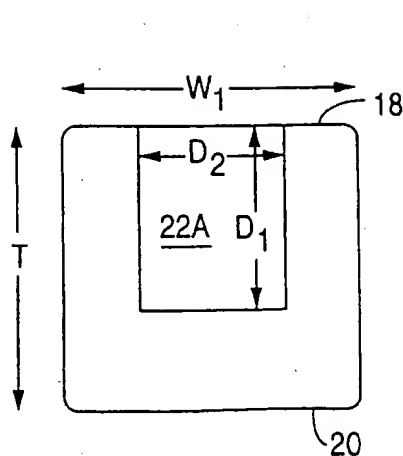


Fig. 3A

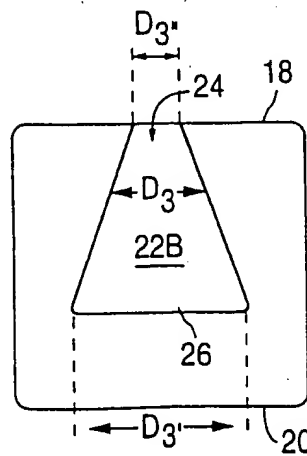


Fig. 3B

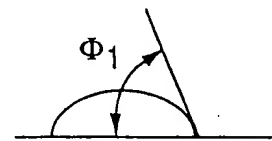


Fig. 4A

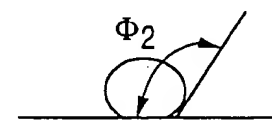


Fig. 4B

INTERNATIONAL SEARCH REPORT

International Application No

PCT/US 00/23535

A. CLASSIFICATION OF SUBJECT MATTER
IPC 7 A61L31/14 A61L31/16

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 A61L A61F

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

EPO-Internal, WPI Data, PAJ, FSTA, INSPEC, COMPENDEX

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO 99 16386 A (SCIMED LIFE SYSTEMS INC) 8 April 1999 (1999-04-08)	1,2, 5-10, 13-15, 30,31, 34-36, 47-59, 63,66-69
Y	page 2, line 20 -page 4, line 11 page 5, line 13 -page 6, line 17 figure 5	1,7, 16-20, 23,24, 27-29, 47,60-65

☒ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

* Special categories of cited documents:

- *A* document defining the general state of the art which is not considered to be of particular relevance
- *E* earlier document but published on or after the international filing date
- *L* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- *O* document referring to an oral disclosure, use, exhibition or other means
- *P* document published prior to the international filing date but later than the priority date claimed

- *T* later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
- *X* document of particular relevance: the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
- *Y* document of particular relevance: the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.
- *Z* document member of the same patent family

Date of the actual completion of the international search

21 December 2000

Date of mailing of the international search report

02/01/2001

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
NL - 2280 HV Rijswijk
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,
Fax: (+31-70) 340-3016

Authorized officer

Menidjel, R

INTERNATIONAL SEARCH REPORT

Internal Application No

PCT/US 00/23535

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	<p>EP 0 875 218 A (ADVANCED CARDIOVASCULAR SYSTEM) 4 November 1998 (1998-11-04)</p> <p>column 5, line 19 - line 50 column 6, line 45 - line 55 column 11, line 10 - column 12, line 25</p>	<p>1,7, 16-20, 23,24, 27-29, 47,60-65</p>
A	<p>EP 0 850 651 A (SCHNEIDER USA INC) 1 July 1998 (1998-07-01)</p> <p>abstract column 1, line 42 - column 2, line 20 column 2, line 45 - column 3, line 16 claims 1-10</p>	<p>1,3,4,7, 11,12, 16-22, 25,26, 30,32, 34,37-46</p>
P,X	<p>PATENT ABSTRACTS OF JAPAN vol. 2000, no. 02, 29 February 2000 (2000-02-29) & JP 11 299901 A (JOHNSON & JOHNSON MEDICAL KK), 2 November 1999 (1999-11-02) abstract</p>	<p>47-63</p>

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/US 00/23535

Patent document cited in search report		Publication date	Patent family member(s)	Publication date
WO 9916386	A	08-04-1999	US 5972027 A	26-10-1999
EP 0875218	A	04-11-1998	US 5843172 A	01-12-1998
			CA 2235031 A	15-10-1998
			JP 10295823 A	10-11-1998
EP 0850651	A	01-07-1998	CA 2225096 A	20-06-1998
			JP 10192413 A	28-07-1998
			US 5980972 A	09-11-1999
JP 11299901	A	02-11-1999	NONE	